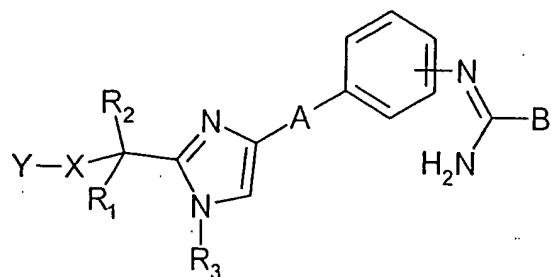


In the claims:

Claim 1 (currently amended)

A compound of the formula



(I)

in the form of racemic, enantiomeric mixture of any combination of these forms,

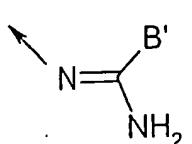
wherein

R₁ is selected from the group consisting of a hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, aryl and aralkyl, the aromatic ring of which is optionally substituted from one to 3 times by a member chosen independently selected from the group consisting of halogen, alkyl and alkoxy;

R₂ is selected from the group consisting of hydrogen, and alkyl and aralkyl;

X is selected from the group ~~consisting~~ consisting of a bond, alkylene of 1 to 5 carbon atoms;

Y is selected from the group consisting of hydrogen, cycloalkyl, NR₄R₅, OR₁₄, SR₁₅ and



or Y is selected from the group consisting of aryl optionally substituted from one to 3 times by a member independently selected from the group consisting of halogen, alkyl and alkoxy;

A is selected from the group consisting of a bond or phenylene;

B and B' are independently selected from the group consisting of alkyl, cycloalkyl, -NR₆R₇, -SR₈, carbocyclic aryl and heterocyclic aryl with 5 to 6 ring members having 1 to 4 heteroatoms selected from the group consisting of O, S and N, said carbocyclic and heterocyclic aryl being optionally substituted by one to three members independently selected from the group consisting of alkyl, alkenyl and alkoxy,

R₃ is selected from the group selected from the group consisting of hydrogen, alkyl and aralkyl;

R₄ is selected from the group consisting of hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, -(C(O)R₉, -C(O)OR₉, -C(O)NHR₉, -SO₂R₉ and aralkyl, the aromatic ring of which is optionally substituted one to 3 times by a member independently selected from the group consisting of hydrogen, alkyl and alkoxy, or R₄ is selected from the group consisting of bis-phenylalkyl,

R₅ is selected from the group consisting of hydrogen, alkyl, aryl and aralkyl, or R₄ and R₅ form with the nitrogen atom which carries them a non-aromatic heterocyclic of five to seven ring members having 1 to 2 heteroatoms, the elements for completing the heterocycle being independently selected from the group consisting of -CHR₁₀-, -NR₁₁-, -O- and -S-;

R₆ and R₇ are selected from the group consisting of -a hydrogen, alkyl, alkenyl and alkynyl;

or R₆ is -NO₂ and R₇ is hydrogen,

or R₆ and R₇ form with the nitrogen atom which carries them a non-aromatic heterocyclic with five to six ring members, the elements for completing the heterocycle being independently selected from the group consisting of -CH₂-, -NR₁₂-, -O- and -S-;

R₈ is alkyl of 1 to 6 carbon atoms optionally substituted from one to 3 times by a member independently selected from the group consisting of halogen, -OH, amino, cyano and aryl;

R₉ is selected from the group consisting of alkyl, haloalkyl, cycloalkyl, cycloalkylalkyl, carbocyclic and heterocyclic aralkyl and aryl, the aromatic ring of which is optionally substituted from one to 3 times by a member independently selected from the group consisting of halogen, alkyl and ~~alkioxy~~ alkoxy;

R₁₀ is selected from a group consisting of hydrogen, alkyl and aryl optionally substituted from one to 3 times by a member independently selected from the group consisting of ~~efp~~ consisting of halogen, alkyl and alkoxy,

R₁₁ is selected from the group consisting of hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, -C(O)R₁₃, -C(O)OR₁₃, -SO₂R₁₃, -C(O)NHR₁₃ and the aryl and aralkyl, ~~toe~~ the aromatic ring of which is optionally substituted from one to 3 times by a member independently selected from the group consisting of halogen, alkyl and alkoxy;

R₁₂ is selected from the group consisting of hydrogen or alkyl;

R₁₃ is selected from the group consisting of alkyl, haloalkyl and carbocyclic and heterocyclic aralkyl or aryl, the aromatic ring of which is optionally substituted one to 3 times by a member independently selected from the group consisting of halogen, alkyl and alkoxy;

R₁₄ is selected from the group consisting of alkyl, phenyl and aralkyl; and

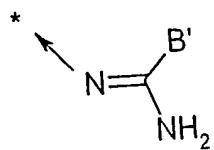
R₁₅ is selected from the group consisting of alkyl, phenyl and aralkyl;

it being understood:

- that alkyl or alkoxy, unless otherwise specified, has 1 to 12 carbon atoms;
- that alkenyl or alkynyl, unless otherwise specified, has has 2 to 6 carbon atoms;
- that cycloalkyl, unless otherwise specified, has 3 to 7 carbon atoms;
- and a salt of a compound of formula (I).

Claim 2 (previously presented) A compound of claim 1, wherein X is a bond or alkylene of 1 to 5 carbon atoms and Y is -NR₄R₅; and a salt thereof.

Claim 3 (currently amended) A compound of claim 1, wherein X is a bond or alkylene of 1 to 5 carbon atoms and Y is



and a salt thereof.

Claim 4 (currently amended) A compound of claim 1, wherein X is a bond or alkylene of 1 to 5 carbon atoms and Y is cycloalkyl or aryl optionally substituted

one to 3 times by a member independently selected from the group consisting of halogen, alkyl and alkoxy; and a salt thereof.

Cancel Claim 5.

Claim 6 (currently amended) A compound of claim 1, selected from the group consisting of

- butyl-2-[4-(4-{{(1Z)-amino(thien-2-yl)methylene]-amino}phenyl}-1*H*-imidazol-2-yl]ethylcarbamate;
- butyl-2-[4-(3-{{(1E)-amino(thien-2-yl)methylene]-amino}phenyl}-1*H*-imidazol-2-yl]ethylcarbamate;
- butyl-2-[4-(4'-{{(1Z)-amino(thien-2-yl)methylene]amino}-1,1'-biphenyl-4-yl}-1*H*-imidazol-2-yl]ethylcarbamate;
- N'-(4-{2-[(cyclohexylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N'-(4-{2-[(cyclohexylamino)ethyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N'-(3-{2-[(cyclohexylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N'-[4-(2-{{cyclohexyl(methyl)amino}methyl}-1*H*-imidazol-4-yl)phenyl]thiophene-2-carboximidamide;
- N'-(4-{2-[(dibenzylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N'-(4-{2-[(benzylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N'-{3-[2-(aminomethyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;

- N -{3-[2-({[(1E)-amino(thien-2-yl)methylene]-amino}methyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{4-[2-({[(1E)-amino(thien-2-yl)methylene]-amino}methyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{3-[2-(2-cyclohexylethyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{3-[2-(1-pentylhexyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{4-[2-(2-cyclohexylethyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{3-[2-(cyclohexylmethyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -{3-[2-(3-cyclohexylpropyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;
- N -[3-(2-hexyl-1*H*-imidazol-4-yl)phenyl]thiophene-2-carboximidamide;
- N -{4-[2-(2-cyclohexylethyl)-1*H*-imidazol-4-yl]phenyl}- N' -nitroguanidine;
- N -{(4-{2-[(cycloheptyl amino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N -(4-{2-[(methylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N -(4-{2-[(cyclobutylamino)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide;
- N -[4-(2-{[(2,2-diphenylethyl)amino]methyl}-1*H*-imidazol-4-yl)phenyl]thiophene-2-carboximidamide;
- N -{3-[2-(2-[(1E)-amino(thien-2-yl)methylene]amino)ethyl]-1*H*-imidazol-4-yl}phenyl}thiophene-2-carboximidamide;
- N -{(3-{2-[(phenylthio)methyl]-1*H*-imidazol-4-yl}phenyl)thiophene-2-carboximidamide and;
- N -{3-[2-(4-isobutylbenzyl)-1*H*-imidazol-4-yl]phenyl}thiophene-2-carboximidamide;

and a salt thereof.

Claim 7 (cancelled)

Claim 8 (currently amended) A pharmaceutical composition containing,
as active ingredient, a compound of claim 1; and an inert pharmaceutical compound
carrier.

Claims 9 and 10 (cancelled)

Claim 11 (currently amended) The method intended to treat or prevent a
disorder/disease selected from the group consisting of treating pain, multiple sclerosis,
disorders of the central or peripheral nervous system, cardiovascular disorders, disorders
of the skeletal muscle and of the neuromuscular joints, inflammatory diseases, hearing
losses of traumatic, acoustic or toxic origin and tinnitus, complications linked with auto-
immune and viral diseases and the neurological diseases associated with intoxication,
treatments and disorders of genetic origin in warm-blooded animals comprising
administering to warm-blooded animals in need thereof an effective amount of a
compound of claim 1.

Cancel Claim 12.